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(56) Documents Cited

EP 0474284 A EP 0189042 A EP 0130647 A
US 6106700 A US 5482689 A US 4108782 A

(58) Field of Search

UK CL (Edition R) E1F FJF FPA FPC FPD
INT CL⁷ E21B
Online: WPI, EPODOC, JAPIO

(54) Abstract Title

Treating subterranean formations using a non-ionic surfactant gelling agent

(57) A subterranean formation is treated, e.g. fractured, acidized, gravel packed, stimulated or fluid loss controlled, by injecting an aqueous, viscoelastic fluid containing a non-ionic amine oxide surfactant gelling agent. The preferred amine oxide is tallow amido propylamine oxide.

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Fig. 1
Surfactant Gel Viscosity In 3% KCl Brine

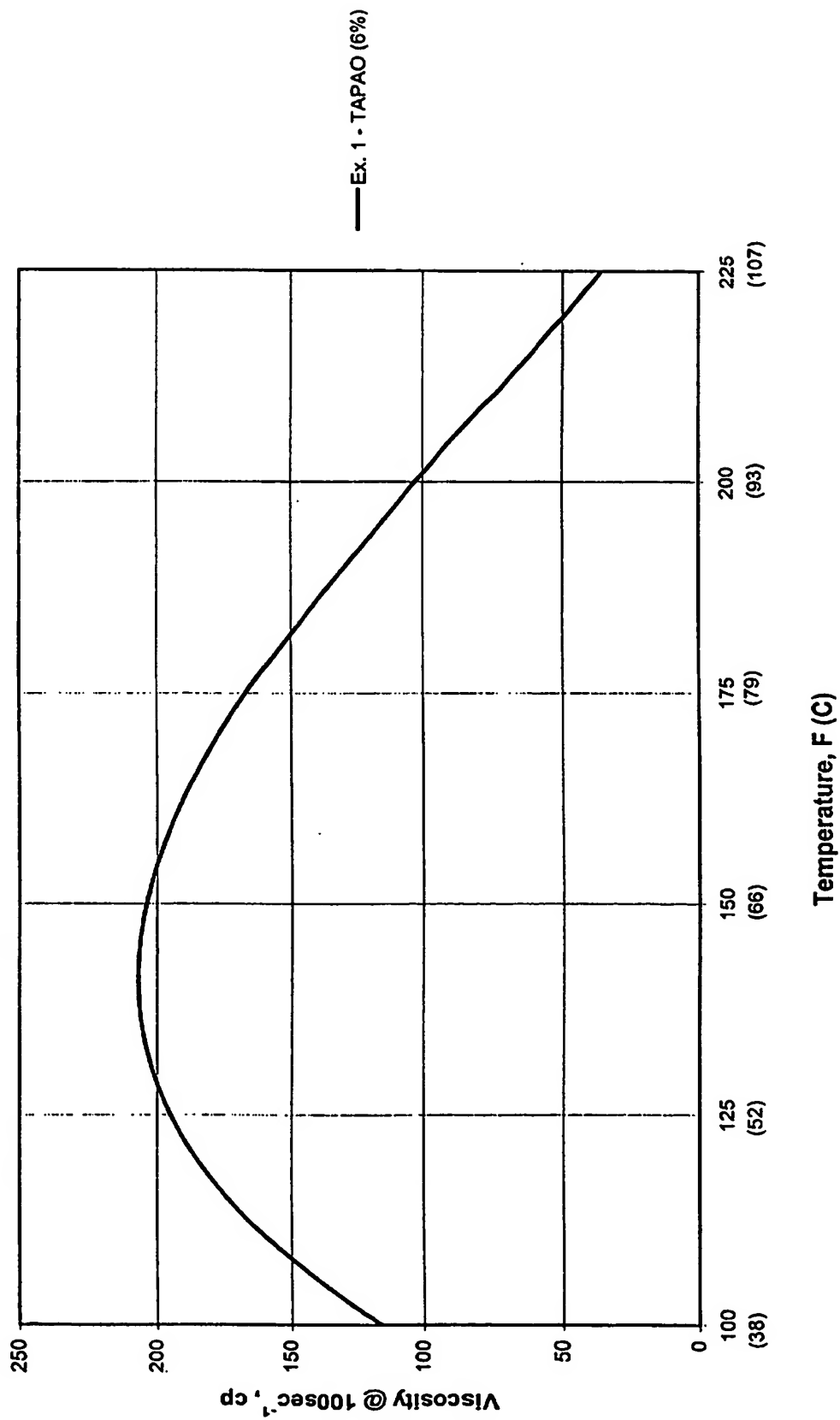
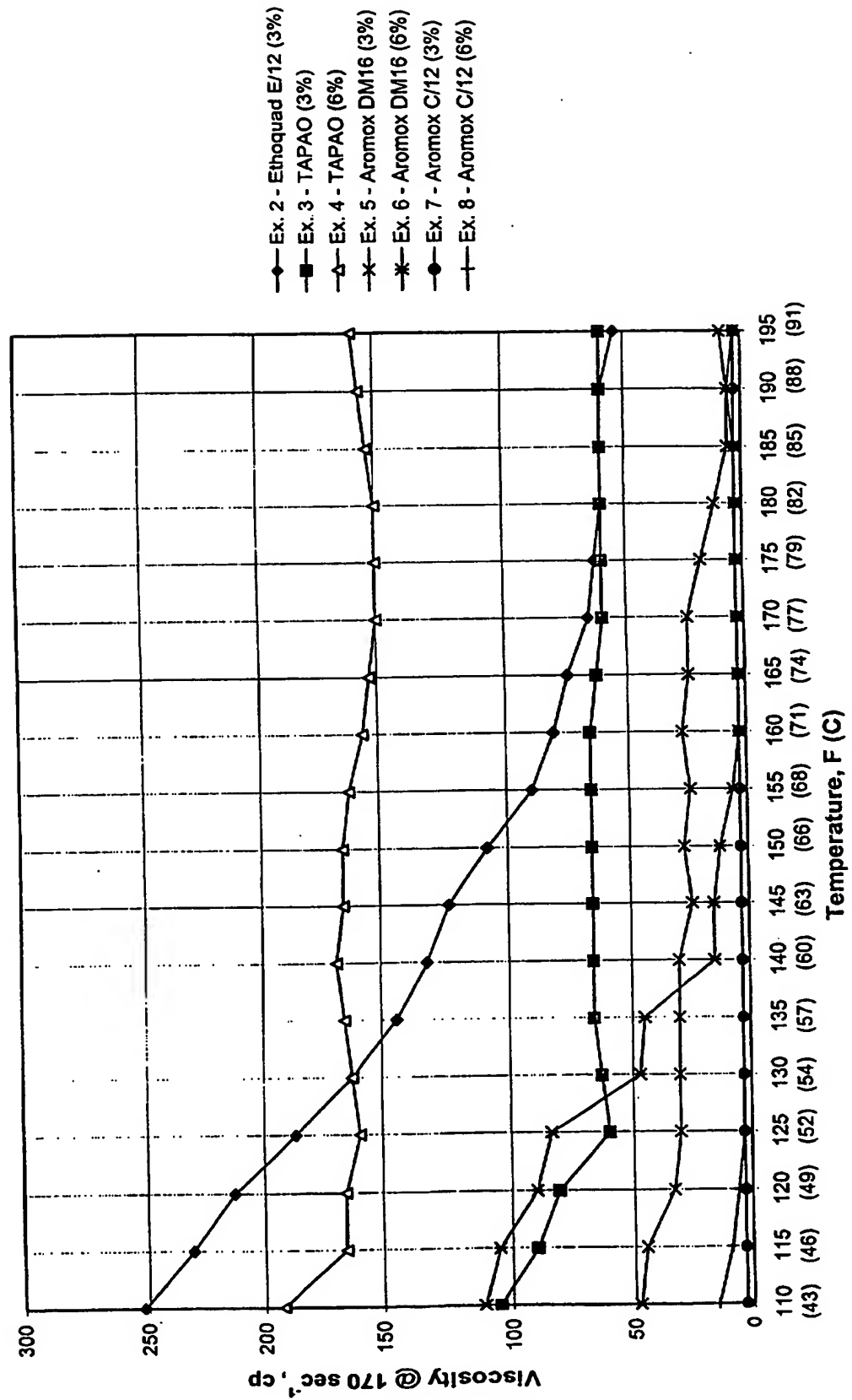


Fig. 2
Viscosities of 3% KCl brine on Fann 35 from 110 to 195F



HYDRAULIC FRACTURING USING NON-IONIC SURFACTANT GELLING AGENT

Field of the Invention

5 The present invention relates to treatment fluids used during petroleum recovery operations, and more particularly relates, in one embodiment, to methods of using treatment fluids containing gelling agents during petroleum recovery operations.

Background of the Invention

10 Hydraulic fracturing is a method of using pump rate and hydraulic pressure to fracture or crack a subterranean formation. Once the crack or cracks are made, high permeability proppant, relative to the formation permeability, is pumped into the fracture to prop open the crack. When the applied pump rates and pressures are reduced or removed from the formation, the crack or fracture cannot close or heal completely
15 because the high permeability proppant keeps the crack open. The propped crack or fracture provides a high permeability path connecting the producing wellbore to a larger formation area to enhance the production of hydrocarbons.

 The development of suitable fracturing fluids is a complex art because the fluids must simultaneously meet a number of conditions. For example, they must be stable at
20 high temperatures and/or high pump rates and shear rates which can cause the fluids to degrade and prematurely settle out the proppant before the fracturing operation is complete. Various fluids have been developed, but most commercially used fracturing fluids are aqueous based liquids which have either been gelled or foamed. When the fluids are gelled, typically a polymeric gelling agent, such as a solvatable
25 polysaccharide is used. The thickened or gelled fluid helps keep the proppants within the fluid.

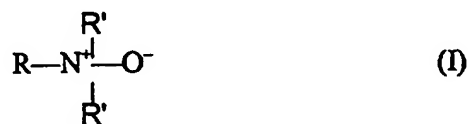
 While polymers have been used in the past as gelling agents in fracturing fluids to carry or suspend solid particles in the brine, such polymers require separate breaker compositions to be injected to reduce the viscosity. Further, such polymers tend to
30 leave a coating on the proppant even after the gelled fluid is broken, which coating may interfere with the functioning of the proppant. Studies have also shown that "fish-eyes" and/or "microgels" present in some polymer gelled carrier fluids will plug pore throats, leading to impaired leakoff and causing formation damage.

 Conventional polymers are also either cationic or anionic which present the
35 disadvantage of likely damage to the producing formations.

anionic gelling agents. The amine oxide technology of this invention has the potential to offer more gelling power per pound, making it less expensive than other fluids of this type.

The amine oxide gelling agents of the invention have the following structure (I):

5



- 10 Where R is an alkyl or alkylamido group averaging from about 8 to 24 carbon atoms and R' are independently alkyl groups averaging from about 1 to 6 carbon atoms. Preferably, R is an alkyl or alkylamido group averaging from about 8 to 16 carbon atoms and R' are independently alkyl groups averaging from about 2 to 3 carbon atoms. A particularly preferred amine oxide gelling agent is tallow amido propylamine oxide
15 (TAPAO), which should be understood as a dipropylamine oxide since both R' groups are propyl.

- The amine oxide gelling agents of the invention may be used in aqueous treatment fluids, particularly brines. The brine base fluid may be any brine, conventional or to be developed which serves as a suitable media for the various
20 concentrate components. As a matter of convenience, the brine base fluid may be the brine available at the site used in the completion fluid, for a non-limiting example.

- While the amine oxide gelling agents of the invention are described most specifically herein as having use in fracturing fluids, it is expected that they will find utility in acidizing fluids, gravel pack fluids, stimulation fluids and the like. Of course,
25 when the treatment fluid is a fracturing fluids, the fluids also contain at least an effective amount of a proppant to prop open the fractures, and the fluid is injected into the formation under sufficient and effective hydraulic pressure and pump rate to fracture the formation. When the treatment fluid is an acidizing fluid, it further contains an effective amount of an acid, either inorganic or organic, of sufficient strength to
30 acidize the formation. When the amine oxide gelling agents are used in a gravel packing fluid, the gelling agent helps contain an effective amount of the gravel within the fluid. If the amine oxide gelling agents are used in another well stimulation fluid, an effective amount of any additional stimulating agent is employed. When the amine oxide gelling agents are used in a fluid loss control application, an effective amount of a
35 salt or easily removed solid is employed, and the amine oxide gelling agents help suspend the salts or solids in the fluid. These other components of the treatment fluids are well known in the art.

5 The effective proportion of the amine oxide gelling agents in the treatment fluids of this invention range from about 0.5 to about 25 vol. %, preferably from about 1 to about 10 vol. %, and most preferably about 6 vol. %. In a non-limiting example, a 6 vol.% solution of the gelling agent is mixed with brine, which is then blended with sand or other particulate, and pumped into a hydrocarbon bearing reservoir.

10 In one non-limiting embodiment of the invention, the non-ionic amine oxide gelling agents are the only gelling agents employed, although more than one may be used. In another non-limiting embodiment of the invention, the non-ionic amine oxide gelling agents are employed in the absence of polymeric gelling agents. In still another non-limiting embodiment of the invention, the non-ionic amine oxide gelling agents are employed in the absence of either cationic or anionic gelling agents.

15 In the method of this invention, breaking the gel of the aqueous viscoelastic treating fluid made using the amine oxides of this invention may be accomplished by a variety of mechanisms. These may include, but are not necessarily limited to, contacting the fluid with a hydrocarbon, contacting the fluid with alkoxylated alcohol solvents, dilution, such as with larger quantities of brine or water, or the addition of a reactive agent. The hydrocarbon may be the hydrocarbon produced from the formation or other hydrocarbon.

20 In another embodiment of the invention, the treatment fluid may contain viscosifying agents, other surfactants, clay stabilization additives, scale dissolvers, biopolymer degradation additives, and other common components.

25 The proppant, solid particle or gravel may be any solid particulate matter suitable for its intended purpose, for example as a screen or proppant, etc. Suitable materials include, but are not necessarily limited to sand, sintered bauxite, sized calcium carbonate, sized salts, ceramic beads, and the like, and combinations thereof. These solids may also be used in a fluid loss control application.

30 A basic method is to inject the proppant into a carrier fluid or treatment brine downstream from the conventional pumps which are delivering the gravel packing fluid, e.g. To do this, the proppant is suspended in the viscosified brine. The proppant may thus be delivered by a small injection pump to the carrier fluid at an injection point downstream from the pumps used to transport the gravel packing fluid or other treatment fluid.

The invention will be further described with respect to the following Examples which are not meant to limit the invention, but rather to further illustrate it.

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EXAMPLE 1

The following fluid was prepared in 3% KCl brine: 6 vol. % TAPAO. The surfactant gel viscosity of the fluids were measured on a Brookfield PVS viscometer at 100 sec⁻¹. The results are plotted on the chart of FIG. 1. It was surprisingly discovered that the viscosity of the fluids using the inventive gelling agents herein remains generally stable over the tested temperature range. It was also surprisingly discovered that the viscosity of the fluids using the inventive gelling agents herein remains generally stable over time as well. Five (5) hours was a typical test period for these tests.

EXAMPLES 2-8

The following fluids were prepared in 3% KCl brine:

Comparative Example 2: 3 vol. % Ethoquad E/12.

Example 3: 3 vol. % TAPAO of a 50 vol. % solution.

Example 4: 6 vol. % TAPAO of a 50 vol. % solution.

Comparative Example 5: 3 vol. % AROMOX DM16

Comparative Example 6: 6 vol. % AROMOX DM16

Comparative Example 7: 3 vol. % AROMOX C/12

Comparative Example 8: 6 vol. % AROMOX C/12

The AROMOX materials are polymeric quaternary ammonium halide salt gelling agents commercially available from Akzo-Nobel, Inc. AROMOX DM16 is a polymeric quaternary ammonium halide salt gelling agent have a C₁₆ substituent and two C₁ substituents on the nitrogen. AROMOX C/12 is a polymeric quaternary ammonium halide salt gelling agent have a C₁₂ substituent and two C₁ substituents on the nitrogen.

The surfactant gel viscosity of the fluids were measured on a Fann 35 viscometer at 170 sec⁻¹. The results are plotted on the chart of FIG. 2. It can be seen again that the fluid of comparative Example 2 using Ethoquad E/12 loses viscosity as the temperature increases. It was again shown that the viscosity of the fluids using the inventive gelling agents herein remains generally stable over the tested temperature range. The viscosity of the fluids using the inventive gelling agents herein (Examples 3 and 4) was also higher and more stable than the comparative Examples 5-8 using commercially available AROMOX materials.

AROMOX E/12 and 50/50 mixtures of AROMOX C/12 with AROMOX E/12 at both 3 vol.% and 6 vol.% were also tested, but gave generally lower viscosities than AROMOX 16 at 3 vol.%.

The inventive non-ionic, non-polymeric amine oxide gelling agents of this invention provide gelling stability over a wide temperature range and at relatively high

temperatures. They are also expected to be relatively non-damaging to the formation since they are non-ionic.

5 In the foregoing specification, the invention has been described with reference to specific embodiments thereof, and has been demonstrated as effective in providing a treatment fluid with stable surfactant gel viscosity. However, it will be evident that various modifications and changes can be made thereto without departing from the broader spirit or scope of the invention as set forth in the appended claims. Accordingly, the specification is to be regarded in an illustrative rather than a restrictive sense. For example, specific combinations of brines, amine oxides and other
10 components falling within the claimed parameters, but not specifically identified or tried in a particular composition, are anticipated to be within the scope of this invention.

Claims

We Claim:

1. A method for treating a subterranean formation comprising:
providing an aqueous viscoelastic treating fluid having:
an aqueous base fluid; and
a non-ionic amine oxide surfactant gelling agent;
injecting the aqueous viscoelastic surfactant treating fluid through a wellbore
and into the subterranean formation; and
treating the subterranean formation under conditions effective to do so.
2. The method of claim 1 where the non-ionic amine oxide surfactant gelling agent is the only gelling agent employed.
3. The method of claim 1 where the non-ionic amine oxide surfactant gelling agent is employed in the absence of a polymeric gelling agent.
4. The method of claim 1 where the non-ionic amine oxide surfactant gelling agent is present in the aqueous base fluid in a proportion from about 0.5 to about 25 vol. %.
5. The method of claim 1 where the non-ionic amine oxide surfactant gelling agent has the formula:



where R is an alkyl or alkylamido group averaging from about 8 to 24 carbon atoms and R' are independently alkyl groups averaging from about 1 to 6 carbon atoms.

6. The method of claim 1 where the non-ionic amine oxide surfactant gelling agent is tallow amido propylamine oxide (TAPAO).
7. The method of claim 1 where the aqueous base fluid is brine.

8. The method of claim 1 where treating the subterranean formation is selected from the group consisting of
- fracturing the formation under effective pressure where the aqueous viscoelastic treating fluid further comprises a proppant;
 - acidizing the formation where the aqueous viscoelastic treating fluid further comprises an acid;
 - packing the formation with gravel where the aqueous viscoelastic treating fluid further comprises gravel;
 - stimulating the formation where the aqueous viscoelastic treating fluid further comprises a stimulating agent;
 - controlling fluid loss where the aqueous viscoelastic treating fluid further comprises a salt or easily removed solid; and mixtures thereof.
9. The method of claim 1 further comprising
- breaking the gel of the aqueous viscoelastic treating fluid by a mechanism selected from the group consisting of contact with a hydrocarbon, contact with alkoxylated alcohol solvents, dilution, and contact with at least one reactive agent.
10. An aqueous viscoelastic treating fluid comprising:
- an aqueous base fluid; and
 - a non-ionic amine oxide surfactant gelling agent.
11. The aqueous viscoelastic treating fluid of claim 10 in the absence of another gelling agent.
12. The aqueous viscoelastic treating fluid of claim 10 in the absence of a polymeric gelling agent.
13. The aqueous viscoelastic treating fluid of claim 10 where the non-ionic amine oxide surfactant gelling agent is present in the aqueous base fluid in a proportion from about 0.5 to about 25 vol. %.
14. The aqueous viscoelastic treating fluid of claim 10 where the aqueous base fluid is brine.

15. The aqueous viscoelastic treating fluid of claim 10 where the non-ionic amine oxide surfactant gelling agent has the formula:



where R is an alkyl or alkylamido group averaging from about 8 to 24 carbon atoms and R' are independently alkyl groups averaging from about 1 to 6 carbon atoms.

16. The aqueous viscoelastic treating fluid of claim 10 where the non-ionic amine oxide surfactant gelling agent is tallow amido propylamine oxide (TAPAO).

17. An aqueous viscoelastic treating fluid comprising:
an aqueous base fluid; and
a non-ionic amine oxide surfactant gelling agent having the formula:



where R is an alkyl or alkylamido group averaging from about 8 to 24 carbon atoms and R' are independently alkyl groups averaging from about 1 to 6 carbon atoms, and
where the non-ionic amine oxide surfactant gelling agent is present in the aqueous base fluid in a proportion from about 0.5 to about 25 vol. %.



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Application No: GB 0023212.4
Claims searched: 1-17

Examiner: David Pepper
Date of search: 14 December 2000

Patents Act 1977
Search Report under Section 17

Databases searched:

UK Patent Office collections, including GB, EP, WO & US patent specifications, in:

UK Cl (Ed.R): E1F FPA, FPD, FJF, FPC.

Int Cl (Ed.7): E21B.

Other: Online: WPI, EPODOC, JAPIO.

Documents considered to be relevant:

Category	Identity of document and relevant passage	Relevant to claims
X	EP 0474284 A (Akzo)	1-5,7-15, 17
X	EP 0189042 A (Hoechst)	1-5,7-15, 17
X	EP 0130647 A (Shell Oil Co)	1-5,7-15, 17
X	US 6106700 A (United Lab Inc)	1-5,7-15, 17
X	US 5462689 A (Clorox Co)	1-5,10-13, 15,17
X	US 4108782 A (The Dow Chemical Co)	1-5,7-15, 17

X	Document indicating lack of novelty or inventive step	A	Document indicating technological background and/or state of the art.
Y	Document indicating lack of inventive step if combined with one or more other documents of same category.	P	Document published on or after the declared priority date but before the filing date of this invention.
&	Member of the same patent family	E	Patent document published on or after, but with priority date earlier than, the filing date of this application.